# SURFACE TENSION OF MELTS OF Na-K-Cs AND Na-K-Rb TERNARY SYSTEMS

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### **ABSTRACT**

In this work the results of the experimental study of surface tension (ST) of alkali metals and their Na-K-Cs and Na-K-Rb ternary systems over the whole composition range and for temperatures from the melting point to 680K are presented. Careful thermal treatment for several hours at 700 K under 10<sup>-7</sup>Pa vacuum condition and a new way of loading the metered cell have been used to exclude possible contamination of the liquid metals and the metered cell. The sessile-drop method, which yields an experimental error of 1% with 95% probability, was used to measure the ST. Contents of impurities were not larger than 0.005%. Surface tension of 120 Na-Cs-K alloys and 48 Na-K-Rb alloys have been measured over the whole composition triangles for the first time. Polytherms of ST for pure metals and ternary alloys are well described by linear equation. The isotherms of ST for each Na/Cs+K section have been constructed. It was found that K is a surface-active addition in ternary alloys when Na:Cs>14:1, while it becomes a surface-inactive component if Na:Cs<14:1. All ternary alloys along the Na:Cs=14:1 section have the same ST, independent from K content. Lines of equal values of ST have been plotted for the whole composition triangles as well. The character of ST data for the Na-K-Rb ternary system is generally similar to that for the Na-K-Cs system.

### INTRODUCTION

Progress of modern science and technology necessitates creation and application of new materials and elaboration of new technological processes. Therefore, alkali metals and their alloys, having a unique combination of such properties as extremely high values of thermal and electric conductivities, the lowest melting temperatures (up to -80°C for some alloys), small values of ionization potential, electronic work function, specific weight and viscosity, high values of surface tension and critical temperature, and so on, attract the investigators' attention. They are widely used in nuclear energetics, new power-intensive chemical current sources, emission electronics, transformers of thermal energy into electric energy, aerospace materials, medicine, and so forth [1-4]. Multicomponent liquid metallic coolants based on alkali metals are a new type of coolants in nuclear energetics.

There are many works in literature devoted to the investigation of physicochemical properties of alkali metals and their alloys [5-9]. However, the surface properties of the latter are not studied well enough due to their high chemical activity. In this work we present a new experimental device and technique for the study of surface properties of

liquid alkali metals. We also present the results of experimental measurements of the ST of pure alkali metals and their ternary alloys.

# DEVICE AND TECHNIQUE FOR THE STUDY OF THE SURFACE TENSION OF LIQUID ALKALI METALS AND THEIR MULTICOMPONENT ALLOYS

For the surface tension measurements the well-known sessile-drop method was used. It is the most precise method. We elaborated a new combined device [10], which allows to form alloys in the entire concentration range and to measure their ST and electronic work function (EWF), all in a single loading. This device permits forming alloys based on either component A or B, of desired concentration and at any stage of the experiments, all without opening the metered cell. The device enables one to measure the ST and EWF of an alloy in identical experimental conditions. The main detail of the device is the metered cell. It was made from molybdenum glass, which does not interact with alkali metals and keeps up ultrahigh vacuum conditions. It does not allow a contact with the external environment, and enables preparation of a large number of alloys with necessary concentrations without unsealing the apparatus. It also allows measuring the ST and EWF for the same alloy many times, renovating the surface of a drop every time. For this technique the density of the alloys must be measured separately. The improved two-capillary pycnometer was used for this purpose. The use of the combined device decreases the duration of measurements by a factor of ten, and yields an essential economy of metals. The measurement errors were 0.5% for the ST, 0.1% for the density, and 1% for EWF. Photoelectric method of Fowler is the most suitable for EWF measurements due to high sensitivity of the results to the state of the surface.

## SURFACE TENSION OF ALKALI METALS AND THEIR TERNARY ALLOYS

Using the combined device and specially elaborated techniques we measured the ST of pure alkali metals and the alloys of the Na-K-Cs and Na-K-Rb systems. The high purity alkali metals with impurity content of no more than 0.005% for Na and Rb and 0.12% for Li were used. The measurements were made over the temperature range 150 to 350°C above the melting point, while both increasing and decreasing the temperature. The drops were formed two or three times for each metal and alloy and were then photographed. Profiles of the drops were measured with a UMM-3 microscope. The polytherms of ST of pure metals over the investigated temperature ranges are described by linear equations:

$$\sigma = \sigma_{\rm m} - a(T - T_{\rm m}), \tag{1}$$

where  $T_m$  is the melting temperature,  $\sigma_m$  is the ST at  $T_m$ , and a is the temperature coefficient of ST. Values of  $T_m$ ,  $\sigma_m$  and a for pure alkali metals are given in Table 1. The results calculated with equation (1) may be recommended as the most reliable.

Table 1. Temperature dependence of surface tension of alkali metals

Metal	Li	Na	K	Rb	Cs
T <sub>m</sub> , K	453.4	370.86	336.71	312.50	301.40
$\sigma_{\rm m}$ , mN/m	419	205.3	116.2	95.8	75.0
-a, mN/m⋅K	0.150	0.094	0.062	0.052	0.050

## SURFACE TENSION OF TERNARY ALLOYS OF ALKALI METALS

No data on the ST of ternary alloys of alkali metals is available in the literature, other than a phase diagram for a single Na-K-Cs [11] system. Beside the combination of the unique properties of pure alkali metals, the ternary alloys have other valuable properties, for example, a wide temperature range of the liquid state. For the Na-K-Cs system it extends from 195 to 2000K (the critical temperature is  $T_c \sim 3000$ K). This property is of great importance due to its application in space nuclear energetics. The experimental results on the ST of 110 alloys of the Na-Cs-K system and 40 alloys of the Na-K-Rb system are presented in this paper for the first time. The alloys of the Na-K-Cs system, formed along ten sections, enclose the entire composition triangle. The alloys were prepared by adding K to a binary Na-Cs alloy at a constant  $X_{\text{Na}}:X_{\text{Cs}}$  ratio. Here  $X_{\text{Na}}$  and  $X_{\text{Cs}}$  are concentrations of Na and Cs.

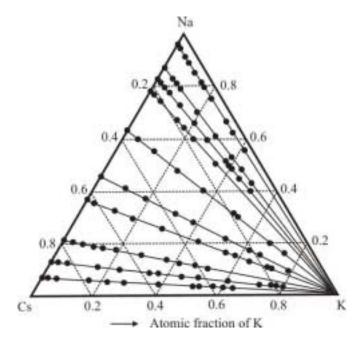


Fig.1. Compositions of 110 ternary alloys of the Na-K-Cs system along the sections, for which the surface tension was measured.

The polytherms of ST of the ternary alloys are described by linear equations with negative or positive temperature coefficients, depending on the concentration of K. The polytherm of the ST of the ternary eutectic alloy is given by equation

$$\sigma(T) = 81.38 - 0.024(T - T_e), \tag{2}$$

where  $T_e=195$  K is the temperature of the ternary eutectic alloy.

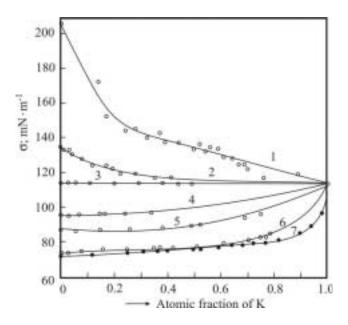


Fig.2. Isotherms of surface tension of ternary alloys of the Na-Cs-K system at 373 K along the sections with  $X_{Na}$ : $X_{Cs} = const$ :

$$1 - \text{Na-K}$$
;  $2 - X_{\text{Na}}$ :  $X_{\text{Cs}}$ =58:1;  $3 - 14$ :1;  $4 - 6$ :1;  $5 - 2$ :1;  $6 - 1$ :2;  $7 - \text{Cs-K}$ .

The isotherms of the ST of ternary alloys at 373 K along five sections with constant  $X_{Na}$ : $X_{Cs}$  ratio are shown in Fig.2. It was found that K is a surface-active component in the Na-K system (Fig.2, curve 1), while it was a surface-inactive component in the Cs-K system (Fig. 2, curve 7). Peculiarity of the influence of K on the ST of Na-Cs-K alloys is in variation of its surface activity when Na is replaced by Cs in the initial Na-Cs alloy. As it is shown in Fig.2, K is a surface-active additive in ternary alloys when  $X_{Na}$ : $X_{Cs}$ >14:1 (curves 1 and 2), and is a surface-inactive additive when  $X_{Na}$ : $X_{Cs}$ <14:1 (curves 4 to 7). This phenomenon may be regarded as a "concentrational inversion of a surface activity" of a component in a ternary system.

Surface tensions of Cs, K and Na are 71, 113.8 and 203.1 mN/m, respectively. The isotherm 3 (Fig.2, curve 3) shows that all ternary alloys along the section beginning with the 93.3 at.% Na + 6.7 at.% Cs binary alloy have the same surface tension of 113.9 mN/m at 373 K, independent of the content of K in the alloy. Independence of the ST of a ternary alloys from the content of one of the components at constant ratio of concentrations of the other

two components is called "concentration buffering of surface tension". The curves of equal values of ST (isolines) of the Na-Cs-K alloys at 373 K are shown in Fig.3. Isoline 6 in Fig.3 is a line of "concentration buffering of ST" at  $X_{Na}$ : $X_{Cs}$ = 14:1.

Forty alloys along four sections of constant  $X_{Na}$ : $X_K$  ratio, formed by addition of Rb, were studied for the Na-K-Rb system. The obtained results on ST are analogous to those for the Na-Cs-K system.

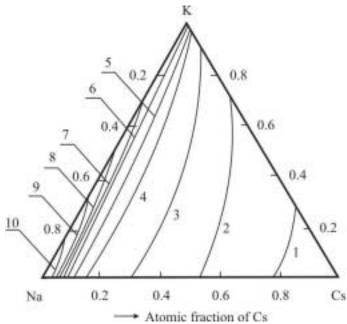


Fig.3. Isolines of surface tension of the Na-Cs-K system at 373 K: 1-75; 2-80; 3-90; 4-100; 5-110; 6-113.8; 7-120; 8-130 mN/m.

# **CONCLUSION**

We conclude with the statement, that interface properties of alkali metals and their alloys depend on the presence of small impurities, the influence of the environment and fields, and the state of a surface. Majority of works devoted to the study of surface properties of liquid alkali metals and alloys, carried out up to 1960s, dealt with metals of insufficient purity, and their results should be considered only as qualitative. Therefore, theoretical and experimental investigation of the influence of gaseous and liquid environments, mechanical and electromagnetic fields, temperature and irradiation of the surface by particles fluxes, adsorption and oxidation processes, and others, on the interface properties are of great importance for managing wetting, spreading and impregnation processes.

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